1675(c)(5)). A record of the Commissioners' votes, the Commission's statement on adequacy, and any individual Commissioner's statements are available from the Office of the Secretary and at the Commission's website.

Participation in the reviews and public service list.—Persons, including industrial users of the subject merchandise and, if the merchandise is sold at the retail level, representative consumer organizations, wishing to participate in these reviews as parties must file an entry of appearance with the Secretary to the Commission, as provided in section 201.11 of the Commission's rules, by 45 days after publication of this notice. A party that filed a notice of appearance following publication of the Commission's notice of institution of the reviews need not file an additional notice of appearance. The Secretary will maintain a public service list containing the names and addresses of all persons, or their representatives, who are parties to the reviews.

For further information concerning the conduct of these reviews and rules of general application, consult the Commission's Rules of Practice and Procedure, part 201, subparts A and B (19 CFR part 201), and part 207, subparts A, D, E, and F (19 CFR part 207).

Please note the Secretary's Office will accept only electronic filings during this time. Filings must be made through the Commission's Electronic Document Information System (EDIS, https://edis.usitc.gov.) No in-person paper-based filings or paper copies of any electronic filings will be accepted until further notice.

Limited disclosure of business proprietary information (BPI) under an administrative protective order (APO) and BPI service list.—Pursuant to section 207.7(a) of the Commission's rules, the Secretary will make BPI gathered in these reviews available to authorized applicants under the APO issued in the reviews, provided that the application is made by 45 days after publication of this notice. Authorized applicants must represent interested parties, as defined by 19 U.S.C. 1677(9), who are parties to the reviews. A party granted access to BPI following publication of the Commission's notice of institution of the reviews need not reapply for such access. A separate service list will be maintained by the Secretary for those parties authorized to receive BPI under the APO.

Staff report.—The prehearing staff report in the reviews will be placed in the nonpublic record on May 2, 2022,

and a public version will be issued thereafter, pursuant to section 207.64 of the Commission's rules.

Hearing.—The Commission will hold a hearing in connection with these reviews beginning at 9:30 a.m. on May 24, 2022. Information about the place and form of the hearing, including about how to participate in and/or view the hearing, will be posted on the Commission's website at https:// www.usitc.gov/calendarpad/ calendar.html. Interested parties should check the Commission's website periodically for updates. Requests to appear at the hearing should be filed in writing with the Secretary to the Commission on or before May 13, 2022. A nonparty who has testimony that may aid the Commission's deliberations may request permission to present a short statement at the hearing. All parties and nonparties desiring to appear at the hearing and make oral presentations should attend a prehearing conference to be held at 9:30 a.m. on May 23, 2022. Oral testimony and written materials to be submitted at the public hearing are governed by sections 201.6(b)(2), 201.13(f), and 207.24 of the Commission's rules. Parties must submit any request to present a portion of their hearing testimony in camera no later than 7 business days prior to the date of the hearing.

Written submissions.—Each party to the reviews may submit a prehearing brief to the Commission. Prehearing briefs must conform with the provisions of section 207.65 of the Commission's rules; the deadline for filing is May 12, 2022. Parties may also file written testimony in connection with their presentation at the hearing, as provided in section 207.24 of the Commission's rules, and posthearing briefs, which must conform with the provisions of section 207.67 of the Commission's rules. The deadline for filing posthearing briefs is June 6, 2022. In addition, any person who has not entered an appearance as a party to the reviews may submit a written statement of information pertinent to the subject of the reviews on or before June 6, 2022. On July 6, 2022, the Commission will make available to parties all information on which they have not had an opportunity to comment. Parties may submit final comments on this information on or before July 11, 2022, but such final comments must not contain new factual information and must otherwise comply with section 207.68 of the Commission's rules. All written submissions must conform with the provisions of section 201.8 of the Commission's rules; any submissions that contain BPI must also conform with

the requirements of sections 201.6, 207.3, and 207.7 of the Commission's rules. The Commission's *Handbook on Filing Procedures*, available on the Commission's website at <a href="https://www.usitc.gov/documents/handbook\_on\_filing\_procedures.pdf">https://www.usitc.gov/documents/handbook\_on\_filing\_procedures.pdf</a>, elaborates upon the Commission's procedures with respect to filings.

Additional written submissions to the Commission, including requests pursuant to section 201.12 of the Commission's rules, shall not be accepted unless good cause is shown for accepting such submissions, or unless the submission is pursuant to a specific request by a Commissioner or Commission staff.

In accordance with sections 201.16(c) and 207.3 of the Commission's rules, each document filed by a party to the reviews must be served on all other parties to the reviews (as identified by either the public or BPI service list), and a certificate of service must be timely filed. The Secretary will not accept a document for filing without a certificate of service.

The Commission has determined that these reviews are extraordinarily complicated and therefore has determined to exercise its authority to extend the review period by up to 90 days pursuant to 19 U.S.C. 1675(c)(5)(B).

Authority: These reviews are being conducted under authority of title VII of the Tariff Act of 1930; this notice is published pursuant to section 207.62 of the Commission's rules.

By order of the Commission. Issued: December 7, 2021.

#### Lisa Barton,

Secretary to the Commission. [FR Doc. 2021–26870 Filed 12–10–21; 8:45 am] BILLING CODE 7020–02–P

# JUDICIAL CONFERENCE OF THE UNITED STATES

## Advisory Committee on Civil Rules; Meeting of the Judicial Conference

**AGENCY:** Judicial Conference of the United States.

**ACTION:** Advisory Committee on Civil Rules; Notice of cancellation of open hearing.

**SUMMARY:** The following virtual public hearing on proposed amendments to the Federal Rules of Civil Procedure has been canceled: Civil Rules Hearing on January 6, 2022. The announcement for this hearing was previously published in the **Federal Register** on August 11, 2021.

DATES: January 6, 2022.

### FOR FURTHER INFORMATION CONTACT:

Bridget Healy, Esq., Acting Chief Counsel, Rules Committee Staff, Administrative Office of the U.S. Courts, Thurgood Marshall Federal Judiciary Building, One Columbus Circle NE, Suite 7–300, Washington, DC 20544, Phone (202) 502–1820, RulesCommittee\_Secretary@ ao.uscourts.gov.

(Authority: 28 U.S.C. 2073.) Dated: December 7, 2021.

#### Shelly L. Cox,

Management Analyst, Rules Committee Staff. [FR Doc. 2021–26868 Filed 12–10–21; 8:45 am]

BILLING CODE 2210-55-P

#### DEPARTMENT OF JUSTICE

## **Drug Enforcement Administration**

[Docket No. DEA-927]

# Importer of Controlled Substances Application: Noramco, Inc.

**AGENCY:** Drug Enforcement Administration, Justice. **ACTION:** Notice of application.

**SUMMARY:** Noramco, Inc., has applied to be registered as an importer of basic class(es) of controlled substance(s). Refer to **SUPPLEMENTARY INFORMATION** listed below for further drug information.

**DATES:** Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before January 12, 2022. Such persons may also file a written request for a hearing on the application on or before January 12, 2022.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. SUPPLEMENTARY INFORMATION: In

accordance with 21 CFR 1301.34(a), this

is notice that on September 22, 2021,

Road, Wilmington, Delaware 19801–4417, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Noramco Inc., 500 Swedes Landing

Controlled substance	Drug code	Schedule
Marihuana	7360 7370 7379 8501 9600 9670	
Noroxymorphone Tapentadol	9668 9780	II   II

The company plans to import Phenylacetone (8501), and Poppy Straw Concentrate (9670) to bulk manufacture other controlled substances for distribution to its customers. The company plans to import an intermediate form of Tapentadol (9780) to bulk manufacture Tapentadol for distribution to its customers. In reference to drug codes 7360 (Marihuana) and 7370 (Tetrahydrocannabinols), the company plans to import a synthetic cannabidiol and a synthetic Tetrahydrocannabinol. No other activity for these drug codes is authorized for this registration.

Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

#### Brian S. Besser,

Acting Assistant Administrator. [FR Doc. 2021–26906 Filed 12–10–21; 8:45 am] BILLING CODE P

## DEPARTMENT OF JUSTICE

# Drug Enforcement Administration [Docket No. DEA-932]

## Bulk Manufacturer of Controlled Substances Application: SpecGX, LLC

**AGENCY:** Drug Enforcement Administration, Justice. **ACTION:** Notice of application.

**SUMMARY:** SpecGX, LLC, has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to Supplemental Information listed below for further drug information.

**DATES:** Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the

issuance of the proposed registration on or before February 11, 2022. Such persons may also file a written request for a hearing on the application on or before February 11, 2022.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

#### SUPPLEMENTARY INFORMATION: In

accordance with 21 CFR 1301.33(a), this is notice that on September 20, 2021, SpecGX LLC, 3600 North 2nd Street, Saint Louis, Missouri 63147, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Phenylacetone	8501	II

The company plans to manufacture the above-listed controlled substance in bulk for conversion to other controlled substances. No other activity for this drug code is authorized for this registration.

#### Brian S. Besser,

Acting Assistant Administrator. [FR Doc. 2021–26907 Filed 12–10–21; 8:45 am] BILLING CODE P

# **DEPARTMENT OF LABOR**

# **Employee Benefits Security Administration**

#### Agency Information Collection Activities; Request for Public Comment

**AGENCY:** Employee Benefits Security Administration (EBSA), Department of Labor.

**ACTION:** Notice.

**SUMMARY:** The Department of Labor (the Department), in accordance with the Paperwork Reduction Act, provides the general public and Federal agencies with an opportunity to comment on proposed and continuing collections of information. This helps the Department assess the impact of its information collection requirements and minimize the public's reporting burden. It also helps the public understand the Department's information collection requirements and provide the requested data in the desired format. The **Employee Benefits Security** Administration (EBSA) is soliciting comments on the proposed extension of